



Drug Enforcement Administration

[Docket No. DEA-932]

Bulk Manufacturer of Controlled Substances Application: SpecGX, LLC.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: SpecGX, LLC, has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application on or before [INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER].

ADDRESS: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on September 20, 2021, SpecGX LLC, 3600 North 2nd Street, Saint Louis, Missouri 63147, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled Substance	Drug Code	Schedule
Phenylacetone	8501	II

The company plans to manufacture the above-listed controlled substance in bulk for conversion to other controlled substances. No other activity for this drug code is authorized for this registration.

Brian S. Besser,
Acting Assistant Administrator.

[FR Doc. 2021-26907 Filed: 12/10/2021 8:45 am; Publication Date: 12/13/2021]